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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/824,851	04/02/2001	Sharat Singh	0225-0033.26	1092

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EXAMINER

TUNG, JOYCE

ART UNIT

PAPER NUMBER

1637

DATE MAILED: 01/03/2003

20

Please find below and/or attached an Office communication concerning this application or proceeding.

# Office Action Summary

Application No.  
**09/824,851**

Applicant(s)  
**Singh et al.**

Examiner  
**Joyce Tung**

Art Unit  
**1637**



-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

## Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136 (a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

## Status

- 1) ☒ Responsive to communication(s) filed on Jul 22, 2002
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11; 453 O.G. 213.

## Disposition of Claims

- 4) ☒ Claim(s) 5-25 is/are pending in the application.
- 4a) Of the above, claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 5-25 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claims \_\_\_\_\_ are subject to restriction and/or election requirement.

## Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on \_\_\_\_\_ is: a) ☐ approved b) ☐ disapproved by the Examiner.  
If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

## Priority under 35 U.S.C. §§ 119 and 120

- 13) ☐ Acknowledgement is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).  
a) ☐ All b) ☐ Some\* c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
  - ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\*See the attached detailed Office action for a list of the certified copies not received.

- 14) ☐ Acknowledgement is made of a claim for domestic priority under 35 U.S.C. § 119(e).  
a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgement is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

## Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892) 4) ☐ Interview Summary (PTO-413) Paper No(s). \_\_\_\_\_
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948) 5) ☐ Notice of Informal Patent Application (PTO-152)
- 3) ☒ Information Disclosure Statement(s) (PTO-1449) Paper No(s). 17 6) ☐ Other:

Art Unit: 1637

***Response to Amendment***

1. The amendment filed 7/22/2002 has been entered.
2. Every rejection in the Office action mailed 1/30/2002 is withdrawn because of the amendment and arguments as filed in the response on 7/22/2002.

**NEW GROUNDS OF REJECTIONS**

***Claim Rejections - 35 USC § 112***

3. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

4. Claims 5-25 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention since the newly added language "an antibody binding compound", "said cleaving agent is a sensitizer and said active species is singlet oxygen or hydrogen peroxide" and "said sensitizer is capable of generating singlet oxygen when photoactivated" has no support in the specification, it constitutes a new matter.

5. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Art Unit: 1637

6. Claims 6-18 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

a. Claims 6-18 are vague and indefinite because of the language "complexes of said electrophoretic probes" has no antecedent basis. Clarification is required.

***Claim Rejections - 35 USC § 103***

7. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103© and potential 35 U.S.C. 102(f) or (g) prior art under 35 U.S.C. 103(a).

8. Claims 5-10,12-14, 16-21 and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Bocuslaski et al. (4,331,590) in view of Giese (Analytical Chemistry, 1983, Vol. 2(7) page 166-168).

Art Unit: 1637

Bocuslaski et al. disclose a specific binding assay involving employing an enzyme-cleaving substrate label in the formation of the labeled conjugate (See column 2, lines 5-9). The labeled conjugates comprise an enzyme-substrate portion, an indicator portion in which the conjugate is cleavable by an enzyme to produce a detectable indicator product (See column 2, lines 5-18) and binding components for antibody (See column 11, lines 24-25). This teaching suggests there is antibody binding compound on the labeled conjugates. There is a linking groups through which the dye indicator is covalently bound to the binding component of the conjugate and cleaving enzyme is to cleave the glycosidic linkage (See column 2, lines 33-36). The assay is adaptable to the detection of any specifically bindable ligand and is particularly useful in the detection of haptens, including antibodies (See column 2, lines 40-43 and column 8, lines 39-62). The label is fluorescence (See column 5, lines 10-39 and column 6, lines 1-10). The bound- and free-species of the labeled conjugate are separated by conventional techniques such as a solid-phase antibody or antigen (See column 10, lines 62-67).

Bocuslaski et al. do not disclose a second antibody binding compound. However, Bocuslaski et al. disclose several labeled conjugates for detecting several different ligands (See column 13, lines 23 to column 18, lines 25). Thus one of ordinary skill in the art would have been motivated to construct a kit including an additional labeled conjugate as needed. In addition Bocuslaski et al. do not disclose the second antibody binding compound having a sensitizer for generating an active species to cleave the cleavable linkage. Since it is unclear what is meant by the phrase "a sensitizer", although the claims are interpreted in light of the

Art Unit: 1637

specification, limitations from the specification are not read into the claims, the teachings of Bocuslaski et al. suggest the limitations of claim 19.

Bocuslaski et al. do not disclose the labeled conjugate which has a mobility modifier as claimed in claim 13, but any molecule compound has mobility and based upon the labeled conjugate, there are 1-500 atoms selected from the group as listed in claims 13 and 19 (See column 5, lines 54-65). This teaching suggests that the labeled conjugate has mobility modifier.

Bocuslaski et al. do not disclose that the labeled conjugate which is released can form distinct peaks upon electrophoretic separation.

Giese disclose an electrophoric release tag which has the same components as the e-tag probe contained in the kit. The release tags comprise 3 molecular groups, known as 'signal', 'release' and 'reactivity' groups. The release group provides a site for specific covalent cleavage and the reactivity group attaches the release tag to a substance of interest (See pg. 166, column 1, first paragraph). Giese also addresses the benefit of using the tag (See pg. 166, column 1, third paragraph to column 2, first paragraph) and the potential usefulness in which the tag can be used for detecting antigen or haptens and the tag can be used for detecting several target simultaneous in a give sample (See pg. 167, column 1, second paragraph). The teachings of Giese suggest that the reactivity group of the tag must have an antibody binding compound as claimed in claim 5.

Bocuslaski et al. and Giese do not disclose a kit which comprising the labeled conjugate or the electrophoric release tag.

Art Unit: 1637

One of ordinary skill in the art would have been motivated to construct a kit at the time of the instant invention including all the reagents needed for detecting the presence or absence one or more target compounds as claimed because constructing a kit including all the elements needed for performing a method was routine practice in the art for convenience at the time of the instant invention and the labeled conjugate or the electrophoretic release tag is very useful in the detection of one or more target compounds as taught by Giese. It would have been prima facie obvious to construct the kit as claimed.

9. Claim 11 and 24-25 are rejected under 35 U.S.C. 103(a) as being unpatentable over Bocuslaski et al. (4,331,590) in view of Giese (Analytical Chemistry, 1983, Vol. 2(7) page 166-168) as applied to claims 5-10, 12-14, 16-21 and 23 above, and further in view of Breslow et al. (6,331,530).

The teachings of Bocuslaski et al. are set forth in section 8 above and Bocuslaski et al. do not address a cleaving agent is a sensitizer which generates an active species, singlet oxygen or said sensitizer which is capable of generating singlet oxygen when photoactivated.

Breslow et al. disclose a linker between two  $\beta$ -cyclodextrin molecules and that a photosensitizer is encapsulated within a matrix, wherein the cleavable linker is cleaved upon exposure to light (See the abstract). Singlet oxygen is produced to cleave the linker (See column 3, lines 47-51).

It have been prima facie obvious to an ordinary skill in the art at the time of instant invention to construct a kit including a cleaving agent which is photosensitizer and can produce

Art Unit: 1637

singlet oxygen to cleave the linker as taught by Breslow et al. because constructing a kit including all the elements needed for performing a method was routine practice in the art for convenience at the time of the instant invention.

10. Claim 15 is rejected under 35 U.S.C. 103(a) as being unpatentable over Bocuslaski et al. (4,331,590) in view of Giese (Analytical Chemistry, 1983, Vol. 2(7) page 166-168) as applied to claims 5-10,12-14, 16-21 and 23 above, and further in view of McGall (5,843,655).

The teachings of Bocuslaski et al. are set forth in section 8 above and Bocuslaski et al. do not address cleavable linkage is cleaved by oxidation and is selected from the group as listed in claim 15

McGall discloses that a disulfide linkage is cleaved by reducing condition, ester linkage is cleaved under basic or nucleophilic conditions (See column 8, lines 16-21) and the released nucleic acid products carry the detectable label (See column 8, lines 16-31).

It would have been prima facie obvious to construct a kit including the probe which has cleavable linkage cleaved by oxidation as taught by McGall because McGall discloses applying a disulfide bond and ester linkage to the probe and that the method is for detecting the presence of cleavable structural feature, such as double stranded nucleic acid with these linkages (See the Abstract) and this teaching suggests that cleavage is efficient. Thus, an ordinary skill in the art would have constructed the kit including the probe with the cleavable linkage as taught by McGall.



Art Unit: 1637

11. Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a).

Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

12. Any inquiries concerning this communication or earlier communications from the examiner should be directed to Joyce Tung whose telephone number is (703) 305-7112. The examiner can normally be reached on Monday-Friday from 8:00 AM-4:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion can be reached at (703) 308-1119 on Monday-Friday from 10:00 AM-6:00 PM.

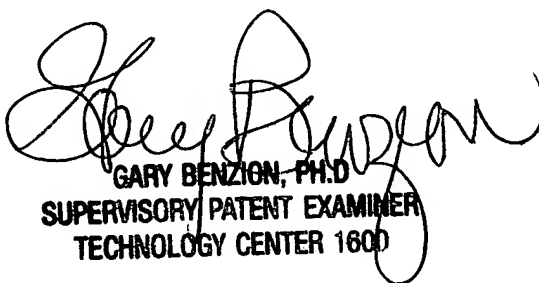
Any inquiries of a general nature or relating to the status of this application should be directed to the Chemical/Matrix receptionist whose telephone number is (703) 308-0196.

Art Unit: 1637

13. Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Art Unit 1637 via the PTO Fax Center located in Crystal Mall 1 using (703) 305-3014 or 308-4242. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989).

Joyce Tung

December 2002

  
GARY BENZION, PH.D.  
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